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Magnetic phase diagram of magnetoelectric LiMnPO₄

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The nature of the spin-flop (SF) transition in the magnetoelectric quasi-2D Heisenberg system LiMnPO₄ is studied in fields applied along the *a* axis. A refinement of the magnetic structure using neutron diffraction data in the SF phase reveals that the spins reorient from being parallel to the *a* axis to be nearly along the *c* axis at magnetic fields between 4 and 4.7 T, depending on temperature. The low-field antiferromagnetic phase boundary is shown to join the spin-flop line tangentially at the so-called *bicritical point*, where there is a suppression of the ordering temperature. At the bicritical field, we observe an increased intensity of the Lorentz broadened elastic scattering at magnetic Bragg peaks above *T_N* as compared to zero field and 10 T, without an increase in peak width. This suggests an increased *density* of fluctuations at the bicritical field as compared to zero field.

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I. INTRODUCTION

Magnetoelectric materials exhibiting a coupling between magnetic and ferroelectric order parameters, such as *RMnO₃*¹ (where *R* is a rare earth element) and the charge-ordered *LuFe₂O₄*,² have drawn particular interests due to scientific challenges in unraveling the coupling mechanism, as well as for their potential applications.^{3,4} The *lithium orthophosphates* *LiMPO₄* (*M* = Mn, Co, Fe, or Ni), which are also candidates for cathode materials,⁵ all exhibit a magnetoelectric (ME) effect in their low-temperature antiferromagnetic (AFM) phases.⁶ Contrary to the other lithium orthophosphates which have only off-diagonal nonzero ME-tensor elements, all the diagonal elements of the ME tensor of LiMnPO₄ are nonzero.⁶ In LiNiPO₄, orbital contributions (in the form of Dzyaloshinsky-Moriya interactions or anisotropic exchange) and detailed knowledge of the magnetic structure are crucial in order to explain the ME effect.⁷ In LiCoPO₄, such orbital contributions are very strong and can also be significant in order to explain the very large ME effect.^{6,8} However, the ground state of the free Mn²⁺ ion has *S* = 5/2 and *L* = 0, and hence the spin-orbit coupling—that also induces the single ion anisotropy term—is practically absent here. In Ref. 9, an investigation of the spin excitations revealed that there are three relatively strong exchange couplings in the *bc* plane (*J_{bc}* ~ 0.5 meV) and two much weaker out-of-plane couplings (*J_a* ~ 0.05 meV) as in the other lithium orthophosphates.^{10,11} As expected, a very weak single ion anisotropy (*D* ~ 0.008 meV) was found, compared to the strongest in-plane nearest-neighbor exchange interactions. Thus, LiMnPO₄ is a pseudo-2D Heisenberg system and has been found to exhibit a spin-flop^{12–15} transition, which is a nearly 90° rotation of antiferromagnetically ordered spins, at 4 T < μ₀*H*_{SF} < 4.7 T applied along the easy *a* axis.^{16,17} The low field magnetic and the spin-flopped structures as determined herein are shown in Fig. 1.

In SF systems, the so-called *spin-flop bicritical point*—where the AFM and SF phases join with the paramagnetic

phase at (*T_b*, *H_b*)—has been extensively investigated, both theoretically^{18–20} and experimentally.^{21,22} In 2D *quantum* Heisenberg systems, spin flopping bears an analogy with the Mott-insulator-to-superconductor transition²³ in high temperature superconductors (where the tuning parameter is the chemical potential instead of the magnetic field), which also show a bicritical point.²⁴ The bicritical SF behavior of a more classical 2D-Heisenberg *S* = 5/2 system has also been studied in this context.²⁵

In this work, we present bulk magnetization and neutron diffraction studies of the SF transition in LiMnPO₄, clarifying the magnetic structure in the SF phase which could be magnetoelectric. We analyze the contribution of the classical dipole-dipole (d-d) to the magnetic anisotropy, and show the necessity of higher order orbital modifications to the otherwise *L* = 0 ground state. In addition, the magnetic phase diagram has been determined for magnetic fields applied along the easy *a* axis, including a thorough investigation of the bicritical point. The shape of the phase boundaries are shown to be in accordance with the prediction of Fisher in Ref. 20. Finally, the neutron diffraction intensity from the critical scattering above *T_N* is shown to be significantly increased near the bicritical point. This is due to the vanishing *effective* anisotropy at the SF field and suggests a coexistence of fluctuations with two different spin polarizations.

II. MAGNETIC STRUCTURE AND ANISOTROPY IN LiMnPO₄

LiMnPO₄ has an orthorhombic crystal structure with space group *Pnma* and lattice parameters *a* = 10.46 Å, *b* = 6.1 Å, and *c* = 4.75 Å.²⁶ It has four magnetic Mn²⁺ ions in each unit cell with spin *S* = 5/2, situated at **r**₁ = (0.278, 0.25, 0.972), **r**₂ = (0.778, 0.25, 0.528), **r**₃ = (0.722, 0.75, 0.028), and **r**₄ = (0.222, 0.75, 0.472), as shown in Fig. 1. At zero field, the system displays long-range AFM order at temperatures

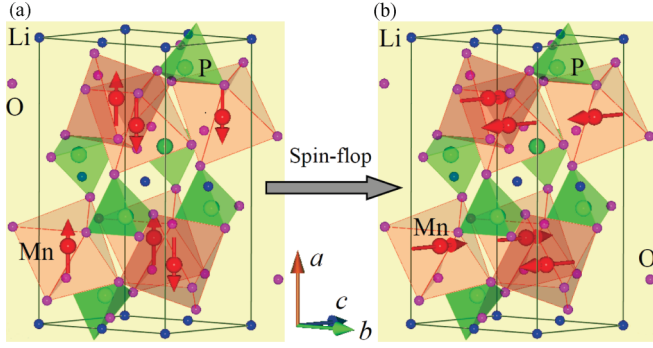


FIG. 1. (Color online) Crystal and magnetic structure of LiMnPO_4 . (a) Zero field magnetic structure polarized along the crystallographic a axis in the AFM phase at low temperatures. (b) The proposed flopped spin structure at $H > H_{\text{SF}}$ applied along a .

below $T_N = 33.85$ K.^{27,28} The magnetic structure can be described in terms of the irreducible representations^{7,29} of the $Pnma$ symmetry group. The zero field AFM spin structure determined in Refs. 27 and 28 has the configuration $(+, +, -, -)$ on the sites \mathbf{r}_i with increasing $i = 1, \dots, 4$ and magnetic moments parallel to the crystallographic a axis, called the C_x -type structure. Basically, this spin structure has ferromagnetic ac planes alternating along b with the periodicity of the lattice, hence described by the wave vector $\mathbf{Q}_{\text{AFM}} = (0, 1, 0)$.

Due to the weak orbital effects ($L = 0$), the magnetic anisotropy of LiMnPO_4 is very small. This implies that the magnetic anisotropy due to the classical dipole-dipole (dd) interaction becomes important. The dd energy can be written as

$$\mathcal{H}_{dd} = -\frac{1}{2} \sum_{ij} \sum_{\alpha\beta} \mathcal{J}_{\alpha\beta}^{dd}(ij) S_i^\alpha S_j^\beta, \quad (1)$$

where i and j denote the magnetic ions in the lattice, and α and β are the $a(x)$, $b(y)$, or $c(z)$ directions in the crystal.³⁰ The sum needs to be over all magnetic ions in the lattice. Using an Ewald's summation method,³¹ the Fourier components of the diagonal dd coupling tensor at \mathbf{Q}_{AFM} can be calculated to be $\mathcal{J}_{aa}^{dd}(\mathbf{Q}_{\text{AFM}}) = 0.0088$ meV, $\mathcal{J}_{bb}^{dd}(\mathbf{Q}_{\text{AFM}}) = -0.0131$ meV and $\mathcal{J}_{cc}^{dd}(\mathbf{Q}_{\text{AFM}}) = 0.0043$ meV.³² The dd interaction contributes to the effective single-ion anisotropy parameters defined in Ref. 9 with contributions which are $D_c^{dd} = 0.00278$ meV and $D_b^{dd} = 0.01370$ meV. The smallest one, i.e., D_c^{dd} , is the one determining the SF field H_{SF} or the spin-wave energy gap at \mathbf{Q}_{AFM} , $\Delta E = g\mu_B\mu_0 H_{\text{SF}}$. Depending on the estimate of the exchange energy (using either the exchange parameters determined by Li *et al.*,⁹ or the magnetization measured above H_{SF} [see Fig. 3(b)]), the value of $\mu_0 H_{\text{SF}} = 4.0$ T, corresponding to $\Delta E = 0.46$ meV, determines the effective value D_c^{eff} to lie in the interval between 0.0050 and 0.0069 meV.³² Hence the dd -interaction is important but accounts only for about half the anisotropy shown by the system. The remaining part has to be due to higher-order orbital modifications of the ground state. These significant orbital modifications are possibly due to the low crystal field symmetry (in LiMnPO_4 , the Mn^{2+} ions are situated within

distorted octahedra), and are expected to increase with decreasing local symmetry. The anisotropy terms induced by these modifications are very weak - but of the same order of magnitude as possible anisotropic exchange terms in LiNiPO_4 $J_{\text{ex}}^{\text{ani}} \sim (\Delta g/g)^2 J_{\text{ex}} \sim 0.01$ meV.^{7,11,33} In LiCoPO_4 however, possible anisotropic exchange terms could be an order of magnitude stronger as $(\Delta g/g)^2 J_{\text{ex}} \sim 0.1$ meV. It should also be noted that the ME effect in LiMnPO_4 is the weakest of the lithium orthophosphates (peaking at 1/40 of that in LiCoPO_4).⁶ Furthermore, these orbital modifications could be the cause of the previously observed zero field weak ferromagnetism in LiMnPO_4 .³⁴

III. EXPERIMENTAL DETAILS

High-quality LiMnPO_4 single crystals were grown by standard flux growth technique.³⁵ The phase purity was confirmed by x-ray powder diffraction. The same single crystal of irregular shape and a weight of 220.14 mg was used in all the experiments presented in this paper.

A CRYOGENIC cryogen free measurement system (CFMS) at the Technical University of Denmark was used to perform vibrating sample magnetization (VSM) measurements. The crystal was oriented on an x-ray Laue camera, glued onto a rigid plastic rod and mounted in the CFMS with the a axis aligned along the magnetic field within 0.4° . Measurements were performed in magnetic fields between 0 and 12 T applied along the crystallographic a axis at $2 \text{ K} < T < 40 \text{ K}$.

To examine the magnetic structure in the SF phase, neutron diffraction experiments were performed using the TriCS diffractometer at the Paul Scherrer Institute (PSI). The crystal was glued to an aluminium sample holder, mounted with the crystallographic a axis vertical within 0.4° in a 6 T Oxford cryomagnet (the maximum field was 4.5 T at the time of the experiment). The cryomagnet has a large vertical opening that allows TriCS to measure three nonequivalent scattering planes. Vertical and horizontal collimations between the sample and the monochromator were both $40'$. The incoming beam propagates in a nose ending in a 10 mm circular aperture. $80'$ collimation was used between the sample and the detector. The neutron wavelength used at TriCS was 1.178 Å.

To examine the magnetic phase boundaries, neutron diffraction experiments were performed using the RITA-II triple axis spectrometer at the PSI with an Oxford 15 T cryomagnet. The incoming and outgoing neutron energy was 5 meV. A $40'$ collimator was inserted between the monochromator and the sample, and a coarse collimation was used between the sample and the analyzer. The crystal was mounted with the b and c axis in the scattering plane. The a axis turned out to be misaligned by about 2° with respect to the vertical field. In the RITA-II experiment, emphasis was on the ordering phase boundaries of the AFM and SF phases. These phase boundaries were measured by performing full omega scans of the (010) peak reflecting both the low field AFM and the SF main structure components, using the same temperature adjustment scheme at all fields. The measurements were all performed when the temperature had stabilized within 0.01° K .

IV. RESULTS AND DISCUSSION

A. Magnetic structure and spin-flop in LiMnPO₄

The *C*-type spin-configuration is mainly reflected in reciprocal lattice points with (even, odd, even) or (odd, even, odd) indices. Furthermore, the magnetic neutron scattering intensity is proportional to the square of the so-called *spin polarization factor* $\mathbf{P}_i(\mathbf{Q}) = \hat{\mathbf{Q}} \times (\hat{\mathbf{e}}_i^m \times \hat{\mathbf{Q}})$, where $i = x, y$, and z is along the crystallographic *a*, *b*, and *c* axes respectively, $\hat{\mathbf{e}}_i^m$ is a unit vector along the spin polarization, and $\hat{\mathbf{Q}}$ is a unit vector directed along the neutron momentum transfer.

In the TriCS experiment 65 reflections were measured at $T = 2$ K and $\mu_0 H = 4.5$ T applied along the *a* axis and in zero field at 60 K in the paramagnetic state. Furthermore, a thorough search was made for zero field spin canting components (as observed in LiNiPO₄⁷) and zero field collinear spin rotations (as observed in LiCoPO₄³⁶) at 2 K, but none were found. A first estimate of the magnetic structure in the SF phase is readily obtained by considering the three key reflections: (010), (012), and (230), listed in Table I. As these peaks mainly reflect a *C*-type structure, but with different spin polarization factors, they will give information about the direction of the magnetic moments in the SF phase, provided that the *C*-type structure remains after the spin flop. This is expected since the spin structure is determined by the exchange constants. The squared spin polarization factors for these three reflections and the observed change in intensity can be found in Table I. The integrated intensity obtained from performing full omega-scans of these reflections as a function of magnetic field is plotted in Fig. 2(a). As evident, the (010) reflection has constant intensity through the SF phase transition within error (error bars represent one standard deviation). This provides evidence that the SF phase retains the zero field *C*-type spin-structure throughout the SF transition and the spin polarization is in the *ac*-plane. The slight increase of the intensity of the (230) and the drastic decrease in the intensity of the (012) peak (see Table I), confirms that the ordered moments are aligned along the *c* axis in the SF phase. The decrease of the (012) peak intensity is less than predicted by the spin polarization factors. However, the crystal size and the magnitudes of the ordered moments ($S = 5/2$) make extinction very probable for the strong reflections. The change of magnetic intensity of the (012) reflection is caused by two simultaneous events: moment rearrangement and extinction. Due to the spin-flop the magnetic intensity decreases and is therefore less affected by extinction. We define the critical SF field at the maximum slope of the (012) peak intensity as a function of field. Figure 2(b) shows the field dependence

TABLE I. Squared spin polarization factors evaluated for spin polarization along *x*, *y* and *z*, for the three peaks used to establish the flopped structure. The expected and observed intensities in the SF phase are given, normalized to the zero field intensity.

	P_x^2	P_y^2	P_z^2	P_z^2/P_x^2	$I_{4.5T}^M/I_{0T}^M$
(0,1,0)	1	0	1	1	1.01(1)
(0,1,2)	1	0.86	0.14	0.14	0.22(1)
(2,3,0)	0.87	0.13	1	1.15	1.10(3)

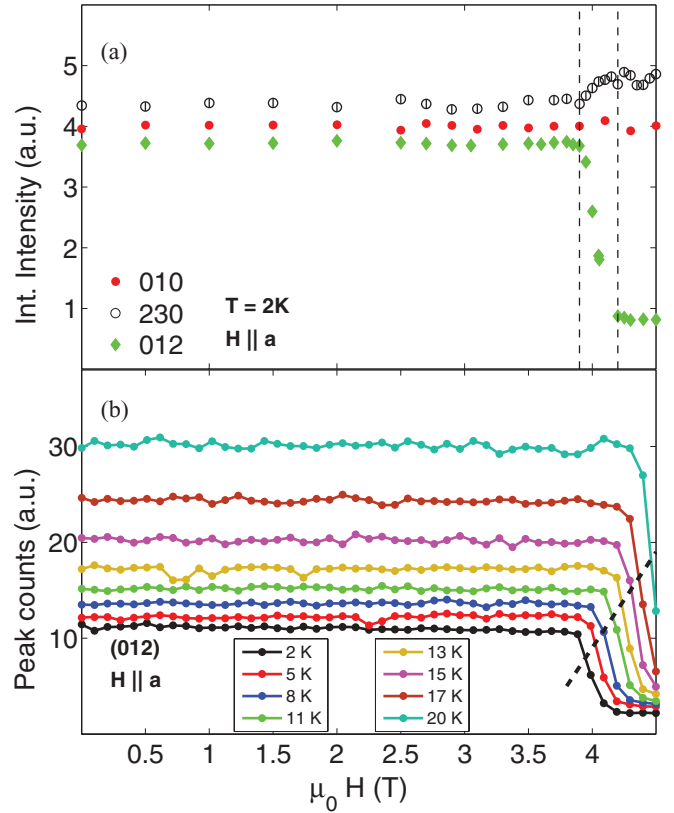


FIG. 2. (Color online) (a) Integrated intensities of the (010), (012), and (230) peaks as a function of field (the intensities of peaks (012) and (230) are scaled by factors of 4 and 6 for comparison). Using Table I, the data are consistent with a *C*-type flop of spin polarization from *a* to *c*. The statistical errors are comparable to or smaller than the markers. (b) Peak intensity of (012) as a function of field at various temperatures (scaled for the sake of illustration) used for phase boundary determination.

of the (012) peak intensity at temperatures between 2 and 20 K. It is evident that the critical field increases slightly with temperature up to 20 K.

A more detailed refinement of the magnetic structure at 2 K and 4.5 T was obtained by using the FULLPROF package.³⁷ Due to the aforementioned extinction effects, the strongest magnetic reflections were not used in the refinements. The remaining 48 measured magnetic peaks were used to refine the structure within a $c_1 C_z + c_2 F_x$ configuration space, where c_1 and c_2 are free parameters and c_2 describes the field induced ferromagnetic component, F_x , along the *a* axis. The result was $c_1 = 3.93(3)\mu_B$ and $c_2 = 0.37(6)\mu_B$ with $\chi^2 = 5.93$, giving a field induced canting angle of $5.4(9)^\circ$ from the *c* axis toward the *a* axis. The relatively high value of χ^2 is most probably due to systematic errors caused by anisotropic absorption in the magnet. To examine the hysteretic behavior of the SF transition, the peak intensity of the (012) reflection was measured at $T = 2$ K for increasing and decreasing field, respectively. The result is plotted in Fig. 3(a) along with the field dependency of the magnetization as obtained from VSM measurements (b) at $T = 2$ K. Neither the neutron diffraction nor the magnetization data presented in Fig. 3 show signs of

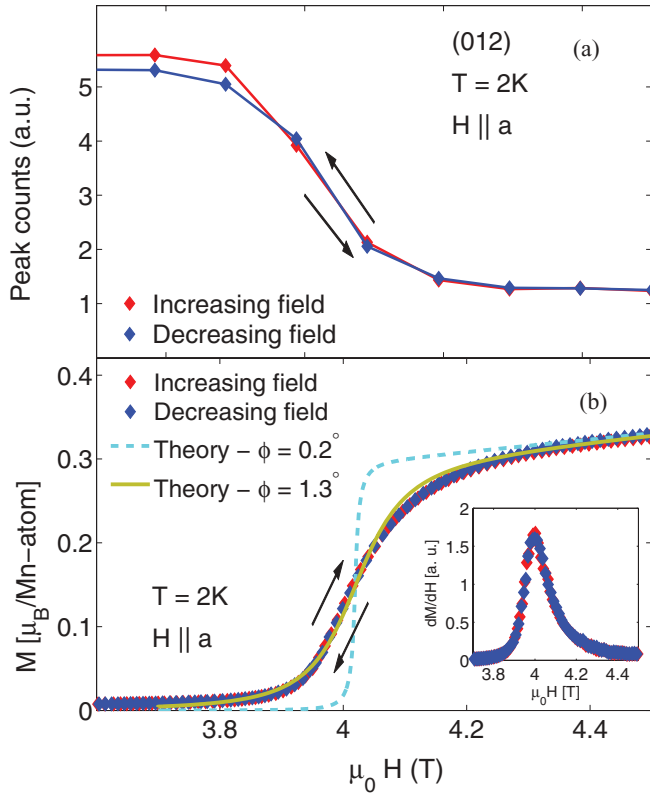


FIG. 3. (Color online) (a) Peak intensity of the (012) peak as a function of applied magnetic field along a at $T = 2$ K. No hysteresis is evident. (b) Magnetization versus field at 2 K. The inset shows the differential magnetization. Neither of them show hysteretic behavior. The field induced moment at $\mu_0 H = 4.5$ T is roughly $0.325\mu_B$ pr. Mn-atom. The critical field is defined as the maximum slope of the curve in both cases. The two theoretical curves represent the mean field calculations described in the text.

hysteresis. It should be noted, however, that an early molecular field analysis of the nature of the spin-flop transition in Ref. 38 predicted that first order behavior can only be expected if the angle between the magnetic field and the easy axis $\phi < \phi_c = 28.6^\circ H_A/H_{EX}$, where H_A and H_{EX} are the anisotropy and exchange fields, respectively. Using the exchange parameters reported in Ref. 9 and the anisotropies determined in this work, $\phi_c \approx 0.2^\circ$ for LiMnPO_4 .³² The magnetization curve at 2 K has been calculated by means of mean field theory using $D_c^{\text{eff}} = 0.0069$ meV, $J(\mathbf{Q}_{\text{AFM}}) - J(0) = 3.15$ meV for field misalignments of $\phi = 0.2^\circ$ and $\phi = 1.3^\circ$, respectively [see Fig. 3(b)].³² The curve for a 1.3° misalignment describes the magnetization data well.

In conclusion, we have shown that the SF structure is confined to the crystallographic ac plane when the field is ramped through the SF transition, and it retains the C -type zero field configuration, but the spins are rotated from the a axis to the c axis with a slight field induced ferromagnetic canting along the field. The field induced moment is roughly $0.325\mu_B$ [as evident in Fig 3(b)] which suggests a canting angle of 3.7° , which is a little lower than the canting angle obtained from the refinement of the neutron diffraction data. A continuous SF transition is observed by both neutron diffraction and VSM measurements. Mean field calculations

account for the magnetization data if the crystal is misaligned by 1.3 degrees with respect to the field. Although the alignment on the Laue camera was within 0.4 degrees, this cannot be assured in the CFMS. Since the SF phase has the same spin configuration as the AFM phase—and as LiNiPO_4 exhibits a ME effect in its C_z -type AFM phase—a magnetoelectric SF phase is indeed possible. A future study of the ME-tensor components in the SF phase of LiMnPO_4 could be helpful in determining the microscopic origin of the ME-effect in this material.

B. Phase diagram and bicritical fluctuations

The spin-flop phase boundary has been measured using both neutron diffraction (TriCS at low temperatures and RITA-II near T_N) and VSM magnetization measurements. The magnetic phase boundary of both the AFM and SF phases has been measured at RITA-II using the (010) peak associated with the C_x and C_z spin components. Figure 4 shows the sub-lattice magnetization $M^+ \propto \sqrt{I_{010}}$, where I_{010} is the integrated intensity of the (010) peak, as a function of temperature at 0 and 4.7 T. The ordering temperature at any given field was found by fitting the intensities—determined by Gaussian fits to the measured rocking curves with a FWHM fixed to the width of the resolution function—to a power-law $M^+(T) = A(T_N - T)^\beta$. The zero field ordering temperature is determined to be $T_N = 33.40(1)$ K, which is a little lower than those determined in Refs. 9 and 27. The critical exponent is determined to be $\beta = 0.150(3)$, which is slightly higher than the previously found value⁹ [$\beta = 0.126(17)$]. This could be due to the fact that in this work only a reduced temperature range in the vicinity around T_N is used for fitting. While a genuine 2D-Heisenberg system does not order, it is well established that weak interplane couplings cause magnetic ordering as observed. The $\beta = 0.150(3)$ value observed in the present system is in full agreement with the result obtained in the classical pseudo-2D system K_2MnF_4 with $\beta = 0.15(1)$.³⁹ Furthermore, there is significantly less critical scattering compared to the results in Ref. 9, where intensity was clearly observed up to $1.5T_N$. However, the energy resolution at RITA-II is much better than that in Ref. 9 (0.2 meV in this

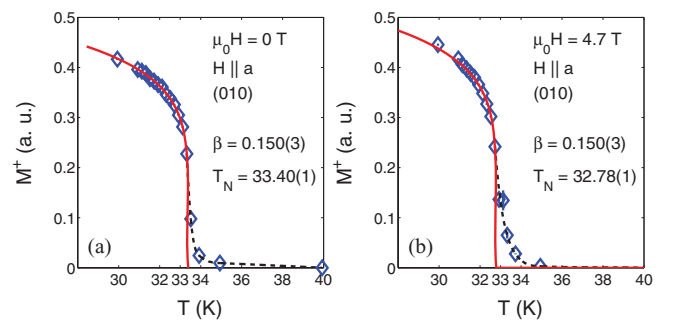


FIG. 4. (Color online) Sublattice magnetization (order parameter) measured by neutron diffraction at RITA-II as a function of temperature in zero field (a), and at $\mu_0 H_b = 4.7$ T (b). The red lines represent power law fits to the staggered magnetization, and the dashed lines are guides to the eye depicting the critical scattering above T_N which is slightly increased at $H = H_b$.

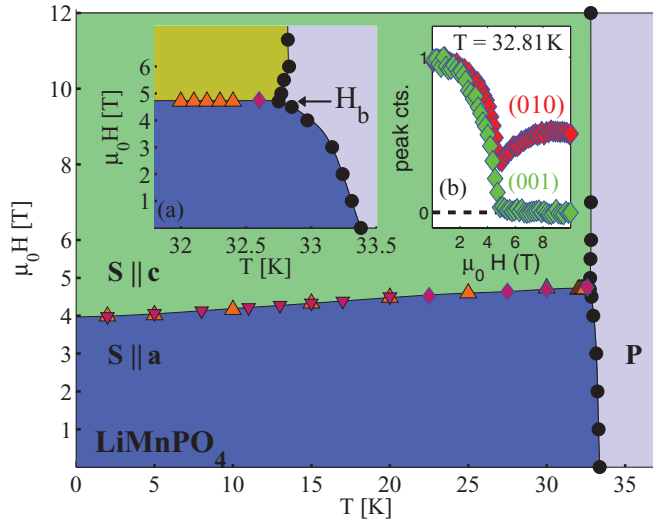


FIG. 5. (Color online) Magnetic phase diagram of LiMnPO_4 for fields applied along a . The up-pointing orange triangles represent the critical fields as measured by VSM magnetization measurements. The down-pointing purple triangles are the critical fields as determined by the TriCS measurements of the (012) reflection, and the purple diamonds represent the critical fields as determined by the RITA-II measurement of the (001) reflection as a function of field. The black circles represent the ordering temperature as found from power-law fits to the integrated intensity of the (010) reflection as measured on RITA-II. Inset (a) shows the phase boundary close to the bicritical point. Inset (b) shows field scans of the (010) and (001) peak intensities at 32.81 K, which is slightly above T_b .

work compared to 1 meV), which supports the suggestion that the critical scattering is dynamical.⁹

Using the same procedure as in zero field, the critical ordering temperature was determined for fields up to 12 T applied along a , with focus on the bicritical point at (T_b, H_b) . The resulting phase diagram is shown in Fig. 5. As evident in Fig. 5 the spin-flop phase boundary is sloping with the critical field increasing from 4 T at 2 K to 4.7 T at the bicritical point. The ordering temperature at the bicritical field

is $T_b = 32.78$ K, which is slightly lower than at zero field. The inset, Fig. 5(a), shows that the AFM phase boundary meets the spin-flop line almost *tangentially*, and the ordering temperatures of both the AFM and SF phases are suppressed at the bicritical field, as predicted by Fisher in Ref. 20. The suppression of the ordering temperature of the SF phase at $\mu_0 H = \mu_0 H_b$ is confirmed by the re-entrance of the (010) intensity above 4.7 T [see the field scan in inset (b) in Fig. 5].

At the bicritical field, two spin polarization states along the a and c directions have the same energy, and the ordering temperature is suppressed. We explore the bicritical region by conducting identical L scans through the magnetic (0 1 0) reflection at various temperatures in the range $T_N + 0.1$ K < $T < T_N + 4$ K, at zero field, at the bicritical field, and at $\mu_0 H = 10$ T. This way the exact same volume of (\mathbf{Q}, ω) space is probed at comparable temperature intervals for the three fields. Figure 6 shows four of these L scans at zero field and at $\mu_0 H_b = 4.7$ T. A significant increase in peak intensity at the bicritical field is evident. We examined whether this increase in amplitude is accompanied by an increase of the correlation length, which proved not to be the case. It should be recalled that with an energy resolution of only 0.2 meV on RITA-II, we measure in the near-elastic regime. However, an increase in the correlation lengths at the bicritical field should still be clearly seen even in this narrow energy interval. The peaks were fit to a Lorentzian [$S \sim 1/(q^2 + \kappa^2)$] that is convoluted with a fixed instrumental Gaussian resolution function. A measure of the correlation lengths along c [$\xi_c(T)$] was found using $\xi = c/2\pi\kappa$, where κ (in r.l.u.) is the half width half maximum of the Lorentzian part of the fit. The results are shown in Fig. 7(a) along with the integrated intensities obtained from the fits [Fig. 7(b)].

The correlation lengths have been fitted to a simple power law [$\xi(T) = A/t(H)^\nu$] where $t(H) = \frac{T - T_N(H)}{T_N(H)}$ is the reduced temperature. The fits are reasonable and similar for all fields, with an average exponent of $\nu = 0.57(6)$. As evident in Fig. 7(a), the three data sets are practically overlapping and there is no significant increase in correlation length at the bicritical field along the crystallographic c axis. Due to the

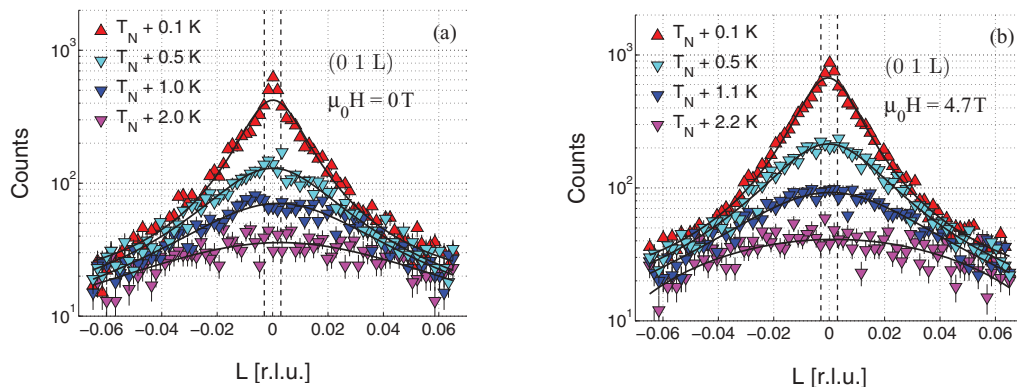


FIG. 6. (Color online) (a) L scans of the (010) peak at four selected temperatures above T_N at $\mu_0 H = 0$ T plotted on a logarithmic scale; the Lorentzian shape is clearly evident. (b) L scans of the (010) peak at similar temperatures relative to T_b at $\mu_0 H = H_b = 4.7$ T. The peak shapes are similar to those obtained at zero field, but with a significantly increased amplitude. The dashed lines indicate the FWHM of the (010) Bragg peak, which is 0.006 r.l.u.

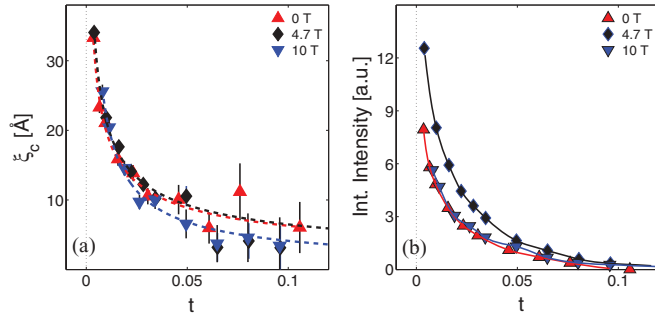


FIG. 7. (Color online) (a) The correlation lengths as a function of temperature relative to $T_N(H)$ as obtained from Voigt fits to peaks such as those depicted in Fig. 6 for 0, 4.7, and 10 T applied along a . The correlation lengths exhibit similar magnitude and dependency on temperature for all fields. The dashed lines mark the fits as described in the text. (b) The integrated intensity as a function of relative temperature for 0, 4.7, and 10 T applied along a . A significant increase in intensity is evident at the bicritical field $\mu_0 H_b = 4.7$ T. The solid lines are guides to the eye.

vertical focusing of the spectrometer, a large part of the Brillouin zone along the vertically oriented a axis is effectively measured. Assuming that there is no change in the correlation length at the bicritical field along the b axis, this suggests that the temperature dependence of the correlation lengths remains unchanged at all fields. Figure 7(b) shows the integrated intensity as a function of t at the three fields. The overall increase of intensity at the bicritical field, as compared with the intensity at 0 and 10 T, is significant. We argue that this increase can only be due to an increased spectral weight $S(\mathbf{Q}, \omega)$ in the measured volume of (\mathbf{Q}, ω) space at the bicritical field. A qualitative interpretation of this increase in critical scattering intensity is that either the *density* or the *lifetime* of the critical fluctuations is increased at the bicritical field. We have no empirical evidence ruling out an increase in the lifetime of the fluctuations. However, due to the field induced isotropy in the ac plane, the most intuitive suggestion is that there is a coexistence of fluctuations in two different spin polarization states, $M_{C_x}^\dagger$ and $M_{C_z}^\dagger$, resulting in an overall increase of the density of critical fluctuations.

V. CONCLUSIONS

In conclusion, the classical d-d interaction has been shown not to be sufficient to explain the weak anisotropy in LiMnPO_4 as obtained from the SF critical field and the spin gap reported in Ref. 9, higher-order orbital modifications to the ground state are necessary. The spin flop in LiMnPO_4 has been determined to occur solely as a reorientation within the ac plane. The SF structure is a C -type configuration polarized along the c axis, consistent with the fact that the b axis is the hard axis of the d-d interaction. The bicritical phase boundaries have been examined and found to be in accordance with theoretical predictions.²⁰ The AFM phase boundary meets the spin flop line almost tangentially, and there is a suppression of the ordering temperatures of both the AFM and SF phase at the bicritical field. Furthermore, the critical fluctuations have been examined at three fields. While there is no significant change in the temperature dependence of the Lorentzian peak widths, we find that the *intensity* of the critical scattering is significantly increased at the bicritical field, both when compared to zero field and to 10 T applied along a . This suggests the coexistence of critical fluctuations of the two different phases with average moment direction along the crystallographic a and c direction, respectively, possibly both magnetoelectric in nature. This increases the overall density of fluctuations near the bicritical point, as observed.

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